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3-1201, Daerim Apt., Sadang-dong, Dongjak-ku, Seoul
156-775 (KR). YI, Jung, Bum [KR/KR]; 113-205,
SuwonHaniltown Apt., 881, Jowon-dong, Jangan-ku,
Suwon-si, Kyunggi-do 440-850 (KR).

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(74) Agent: SUH, Jong, Wan; 7th Fl., New-Seoul Bldg.,
828-8, Yeoksam-dong, Gangnam-ku, Seoul 135-080 (KR).

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(71) Applicant (for all designated States except US): PACIFIC
CORPORATION [KR/KR]; 181, Hankang-ro 2-ka,
Yongsan-ku, Seoul 140-777 (KR).

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(54) Title: NOVEL THIOUREA COMPOUNDS AND THE PHARMACEUTICAL COMPOSITIONS CONTAINING THE SAME

(57) Abstract: The present invention relates to thiourea compounds and the pharmaceutical compositions containing the same, and particularly, to novel thiourea compounds as an antagonist against vanilloid receptor (VR) and the pharmaceutical compositions thereof. As diseases associated with the activity of vanilloid receptor, pain, acute pain, chronic pain, neuropathic pain, post-operative pain, migraine, arthralgia, neuropathies, nerve injury, diabetic neuropathy, neurodegeneration, neurotic skin disorder, stroke, urinary bladder hypersensitivity, irritable bowel syndrome, a respiratory disorder such as asthma or chronic obstructive pulmonary disease, irritation of skin, eye or mucous membrane, farscence, stomach-duodenal ulcer, inflammatory bowel disease and inflammatory diseases can be enumerated. The present invention provides a pharmaceutical composition for prevention or treatment of these diseases.

**Novel thiourea compounds and the pharmaceutical compositions containing the
same**

Technical Field

5 The present invention relates to thiourea compounds and the pharmaceutical compositions containing the same, and particularly, to thiourea compounds with superior efficacy as an antagonist against vanilloid receptor (VR) and the pharmaceutical compositions thereof.

10 **Background Art**

As diseases associated with the activity of vanilloid receptor, pain, acute pain, chronic pain, neuropathic pain, post-operative pain, migraine, arthralgia, neuropathies, nerve injury, diabetic neuropathy, neurodegeneration, neurotic skin disorder, stroke, urinary bladder hypersensitivity, irritable bowel syndrome, a respiratory disorder 15 such as asthma or chronic obstructive pulmonary disease, irritation of skin, eye or mucous membrane, fervesence, stomach-duodenal ulcer, inflammatory bowel disease and inflammatory diseases can be enumerated. The present invention provides pharmaceutical compositions for prevention or treatment of these diseases.

Yet, the diseases described above are only for enumeration, not to limit the scope of clinical application of vanilloid receptor antagonist.

Capsaicin (8-methyl-N-vanillyl-6-nonenamide) is a main pungent component in 5 hot peppers. Hot peppers have been used, for a long time, not only as a spice but also as traditional medicine in the treatment of gastric disorders and when applied locally, for the relief of pain and inflammation (Szallasi and Blumberg, 1999, Pharm, Rev. 51, pp159-211). Capsaicin has a wide spectrum of biological actions, and not only exhibits effects on the cardiovascular and respiratory systems but also induces pain and 10 irritancy on local application. Capsaicin, however, after such induction of pain, induces desensitization, both to capsaicin itself and also to other noxious stimuli to make the pain stopped. Based on this property, capsaicin and its analogues such as olvanil, nuvanil, DA-5018, SDZ-249482, resiniferatoxin are either used as analgesic agent, therapeutic agent for incontinentia urinae or skin disorder, or under development 15 (Wriggleworth and Walpole, 1998, Drugs of the Future 23, pp 531-538).

Transmissions of mechanical, thermal and chemical noxious stimuli are mainly occurred by primary afferent nerve fibers of fine unmyelinated nerve (C-fiber) and thin myelinated nerve (A-fiber), and main reaction site of capsaicin and its analog called

vanilloid is present at the nerve fiber transmitting the noxious stimuli. Capsaicin acts at the receptor existing on these neurons to induce potent stimuli by causing potent inflow of mono-and di-valent cations such as calcium and sodium, then exhibits potent analgesic effect by blocking the nervous function (Wood et al., 1988, J. Neurosci, 8, 5 pp3208-3220). Vanilloid receptor (VR-1) has been recently cloned and its existence becomes clear(Caterina et al., 1997, Nature 389, pp816-824). It was clarified that this receptor transmits not only stimuli by capsaicin analogues(vanilloid) but also various noxious stimuli such as proton and thermal stimuli (Tominaga et al., 1998, Neuron 21, pp531-543). Based on this, it is considered that vanilloid receptor functions as a 10 integrative modulator against various noxious stimuli and carries out critical role in transmissions of pain and noxious stimuli. Recently, knock-out mouse in which gene encoding for vanilloid receptor was deleted was prepared (Caterina et al., 2000, Science 288, pp306-313; Davis et al., 2000, Nature 405, pp183-187). Compared to normal mice, the mouse was found out to exhibit much reduced reaction to thermal stimuli and 15 thermal pain, while exhibiting no difference in general behavior, reconfirming the importance of the receptor in transmission of noxious signal. However, except proton, no other endogenous ligand, not exogenous ligand such as capsaicin, actually involved in transmission of noxious stimuli at vanilloid receptor was known. It is considered that leucotriene metabolite represented by 12-hydroperoxyeicosatetraenoic acid

(12-HPETE) (Hwang et al., 2000, PNAS 11, pp6155-6160) and arachidonic acid derivatives such as anandamide (Zygmunt et al., 2000, Trends Pharmacol. Sci. 21, pp43-44) act as the most likely endogenous ligand for the receptor and proton acts as a cofactor with receptor-stimulating activity, rather than as a direct ligand.

5

As such, a capsaicin-sensitive sensory nerve cell and a vanilloid receptor existing in the cell are distributed over the entire body and play basic function in transmission of noxious stimuli and pain, further act as crucial factor in expression of neurogenic inflammation, thereby to have close relation with the cause of neuropathies, 10 nerve injury, stroke, asthma, chronic obstructive pulmonary diseases, urinary bladder hypersensitivity, irritable bowel syndrome, inflammatory bowel disease, fervescence, skin disorder and inflammatory disease. Lately, their correlation even with neuropathic disease is suggested (WO 99/00125). Recently, attention has focused to the role of afferent sensory nerve responding to capsaicin in gastrointestinal injury, and 15 it was proposed that the afferent nerve might have a dual character that it exhibits protective action against gastric damage by improving gastric microcirculation through releasing peripheral neuropeptide such as CGRP (calcitonin gene-related peptide), while inducing gastric injury by stimulating sympathetic nervous system (Ren et al., 2000, Dig. Dis. Sci. 45, pp830-836). It is determined that vanilloid receptor antagonist has

very high potential to be used for prevention or treatment of the said various diseases by blocking the vanilloid receptor conducting such varied functions.

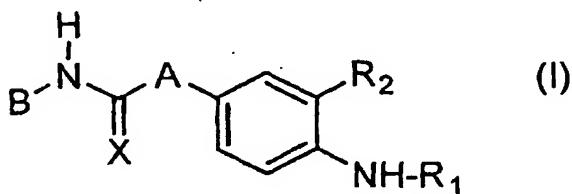
Though it may be, theoretically, anticipated that antagonist for this receptor would exhibit substantial degree of inhibitory action against pain and neurogenic 5 inflammation, it was found out that the competitive antagonist for this receptor, capsazepine, almost the only one known until now, failed to exhibit significant analgesic and anti-inflammatory effects (Perkins and Campbell, 1992, Br. J. Pharmacol. 107, pp329-333). Therefore, not much progress was made on this field. However, recently, there has been a report on significant results for analgesic action of 10 capsazepine in animal studies (Kwak et al., 1998, Neurosci. 86, pp619-626; Santos and calixto, 1997, Neurosci. Lett. 235, pp73-76), in particular, the inventors of the present invention clearly demonstrated through animal studies the analgesic and anti-inflammatory effects of the strong vanilloid receptor antagonists which were identified through experiments in laboratory, and based on this, strongly suggested the 15 development potential of vanilloid receptor antagonist as an analgesic and anti-inflammatory agent. Yet, though the vanilloid receptor antagonist derived from the present studies will mainly act based on the antagonistic activity of itself, even a possibility that it could exhibit the pharmacological activity through transformation into agonist via metabolism after absorption into body is not to be excluded.

To resolve the problems described above, the present invention is to provide novel compounds which are selectively antagonistic to vanilloid receptor and exhibit analgesic and anti-inflammatory effects while causing no irritancy, and pharmaceutical compositions containing the same.

5

Disclosure of the Invention

In order to attain the above objects, the present invention provides a novel compound of formula (I):



10

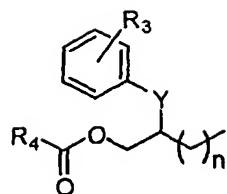
wherein

X represents a sulfur atom or an oxygen atom;

R₁ represents a lower alkyl sulfonyl group having 1 to 5 carbon atoms, an aryl sulfonyl group or a lower alkyl carbonyl group having 1 to 5 carbon atoms, which may be unsubstituted or substituted with halogen atom;

15

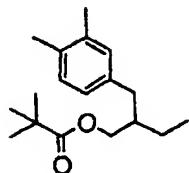
R₂ represents a hydrogen atom, a methoxy group or a halogen;



when A is $-\text{NHCH}_2-$, B represents

(wherein, n is 0

or 1), or when A is $-\text{CH}_2-$, B represents 4-t-butylbenzyl, 3,4-dimethylphenylethyl,



or an oleyl group;

Y represents $-\text{CH}_2-$ or $-\text{CH}_2\text{CH}_2-$;

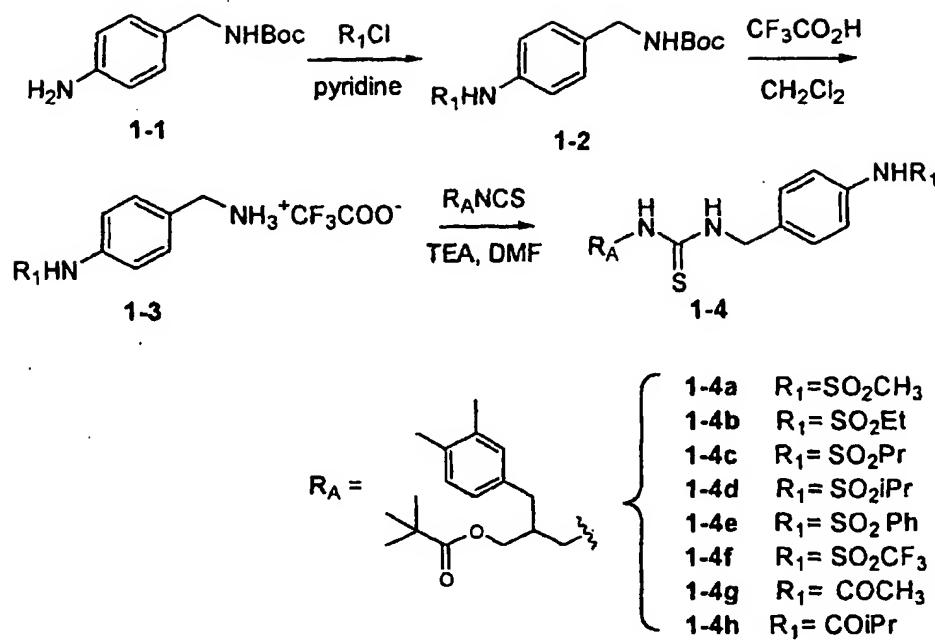
5 R_3 represents hydrogen atom or a lower alkyl group having 1 to 5 carbon atoms; and

R_4 represents a lower alkyl group having 1 to 5 carbon atoms or phenyl group.

The present invention also provides a pharmaceutical composition comprising a 10 compound of formula (I) or a pharmaceutically acceptable salt thereof as an active ingredient.

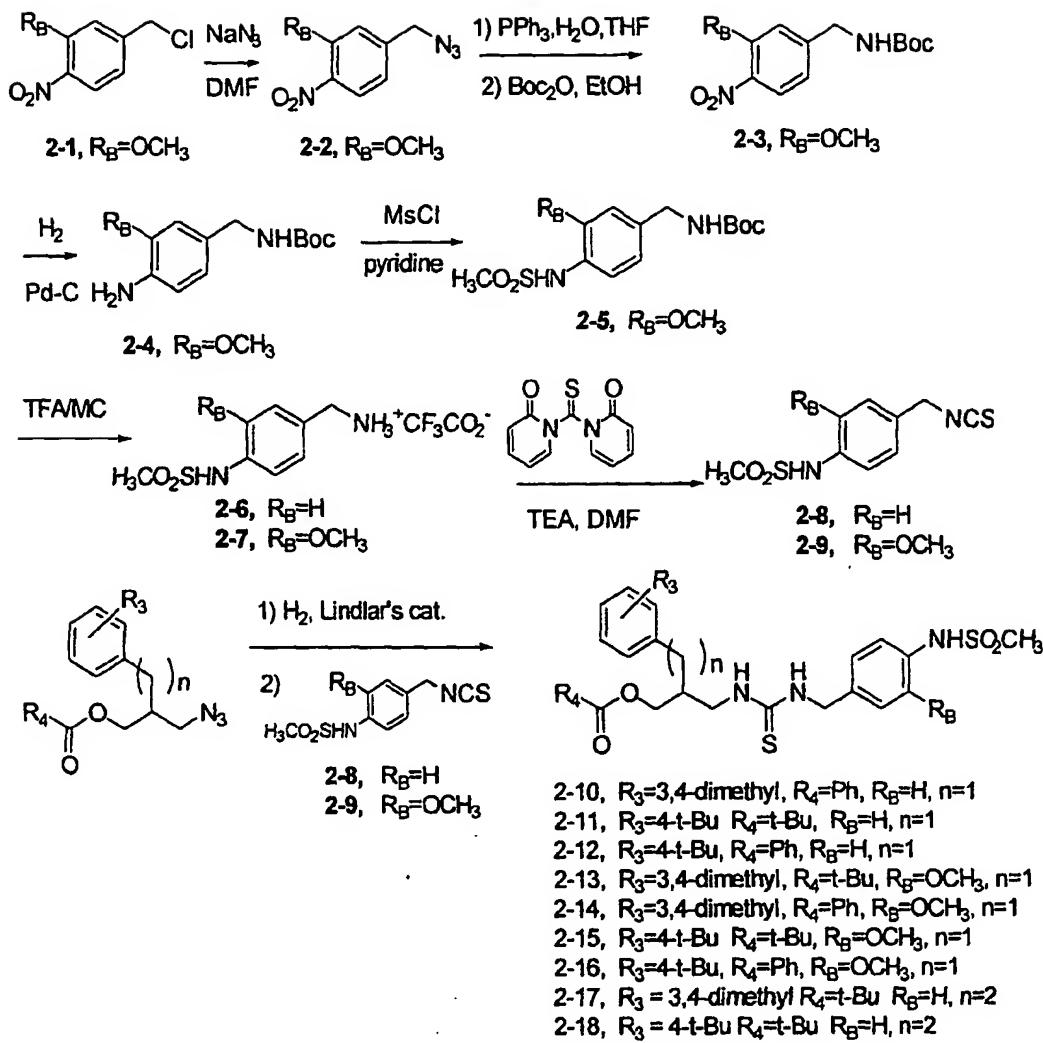
The compounds according to the present invention can chemically be synthesized by the following reaction schemes. However, these are given only for illustration of the invention and are not intended to limit to them.

[SCHEME 1]



5 As depicted in the above Scheme 1, a compound 1-1 was synthesized by selectively protecting 4-aminobenzylamine with Boc(t-butoxycarbonyl) group. Amine of formula 1-1 was reacted with various kinds of agents, e.g., sulfonylchloride or acyl chloride to synthesize a compound 1-2, Boc group was removed from the compound 1-2 under acidic conditions, and then various kinds of isothiocyanate-based compounds 10 were reacted therewith to prepare compounds 1-4a ~ 1-4h.

[SCHEME 2]

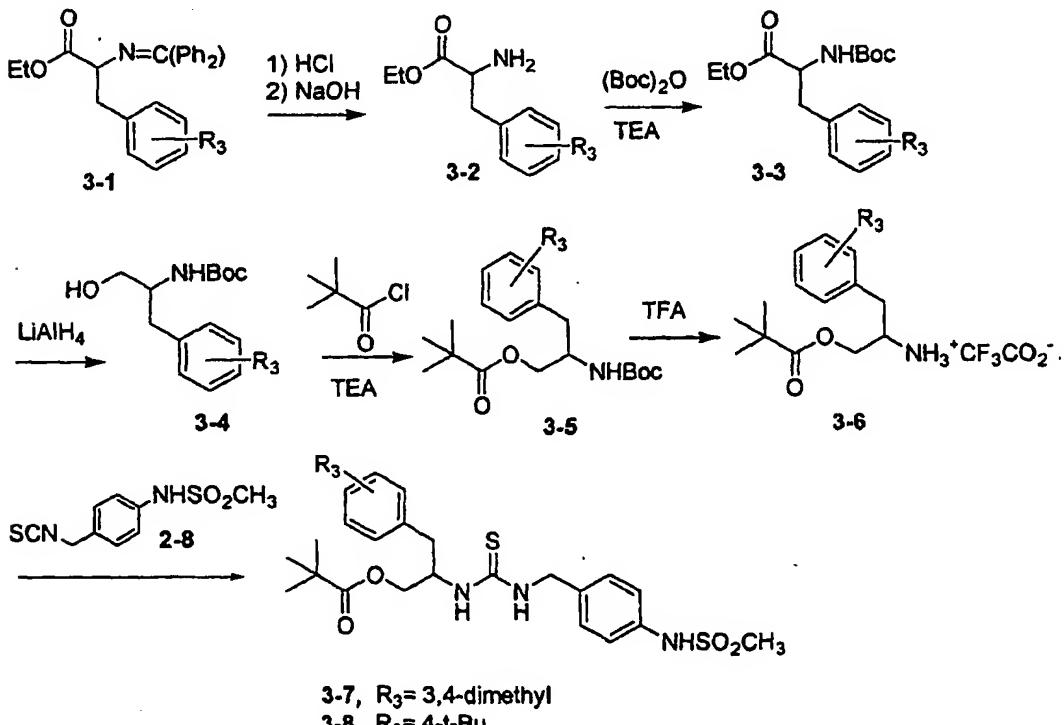


As depicted in the above Scheme 2, azide group was introduced into 3-methoxy-4-nitrobenzyl chloride, reduced, and then protected with Boc group to synthesize a compound 2-3. Nitro group of the compound 2-3 was reduced to produce an amine compound 2-4, methanesulfonylamino group (mesyl group) was introduced thereinto to synthesize a compound 2-5, and then Boc group was removed therefrom to

synthesize an amine compound 2-7. The amine compound 2-7 was treated with 1,1-thio-1H-carbonyl-di-2-pyridone to synthesize an isothiocyanate compound 2-9 as a main intermediate. The azide compounds previously prepared was reacted with the compound 2-9 to prepare compounds 2-13 ~ 2-16.

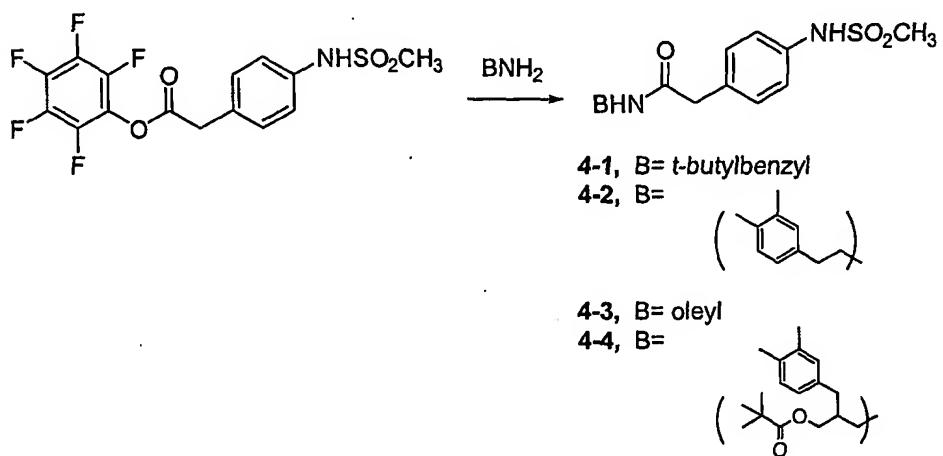
5 In case R_B is H, a compound 2-8 was synthesized in accordance with the above process except that 4-nitrobenzylchloride was used as a starting material, and the compound 2-8 was reacted with azide compound to produce compounds 2-10 ~ 2-12, 2-17 and 2-18.

[SCHEME 3]



As depicted in the above Scheme 3, amine salt 3-6 was synthesized by using a compound 3-1 as a starting material according to a general method, and then the amine salt 3-6 was reacted with an isocyanate compound 2-8 to produce compounds 3-7 and 3-8.

[SCHEME 4]



Compounds 4-1 ~ 4-4 were produced by condensing pentafluorophenyl ester of 4-methanesulfonylaminophenylacetic acid with the known amines.

10

The compound of formula (I) according to the present invention can be provided as a pharmaceutical composition containing pharmaceutically acceptable carriers, adjuvants, or diluents. For instance, the compounds of the present invention

can be dissolved in oils, propylene glycol or other solvents which are commonly used to produce an injection. Suitable examples of the carriers include, physiological saline, polyethylene glycol, ethanol, vegetable oils, isopropyl myristate, etc., but are not limited to them. For topical administration, the compounds of the present invention can be 5 formulated in the form of ointments and creams.

The pharmaceutical composition containing the compound of the present invention as an active ingredient can be used for treating acute, chronic, inflammatory or neuropathic pains; treating urinary bladder hypersensitivity or irritable bowel syndrome (IBS); treating asthma; preventing or treating neurodegenerative diseases; or 10 preventing or treating neurotic skin disorder, or irritation of skin, eye or mucous membrane.

Hereinafter, the formulating methods and kinds of excipients will be described, but the present invention is not limited to them.

15 The compound according to the present invention may also be used in the forms of pharmaceutically acceptable salts thereof, and may be used either alone or in combination or in admixture with other pharmaceutically active compounds.

The compounds of the present invention may be formulated into injections by

dissolving, suspending or emulsifying in water-soluble solvent such as saline and 5% dextrose, or in water-insoluble solvents such as vegetable oils, synthetic fatty acid glyceride, higher fatty acid esters and propylene glycol. The formulations of the invention may include any of conventional additives such as dissolving agents, isotonic agents, suspending agents, emulsifiers, stabilizers and preservatives.

The preferable dose level of the compounds according to the present invention depends upon a variety of factors including the condition and body weight of the patient, severity of the particular disease, dosage form, and route and period of administration, but may appropriately be chosen by those skilled in the art. The compounds of the present invention are preferably administered in an amount ranging from 0.001 to 100 mg/kg of body weight per day, and more preferably from 0.01 to 30 mg/kg of body weight per day. Doses are administered from once to several portions per day. The compounds of the present invention must be present in a pharmaceutical composition in an amount of 0.0001 ~ 10% by weight, and preferably 0.001 ~ 1% by weight, based on the total amount of the composition.

The pharmaceutical composition of the present invention can be administered to a mammalian subject such as rat, mouse, domestic animals, human being and the like via various routes. The methods of administration which may easily be expected

include oral and rectal administration; intravenous, intramuscular, subcutaneous, intrauterine, duramatal and intracerebroventricular injections.

Best Mode for Carrying Out the Invention

5 The present invention is more specifically explained by the following examples. However, it should be understood that the present invention is not limited to these examples in any manner.

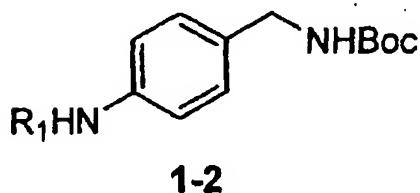
Examples 1 ~ 8

10 The Examples 1~8 are divided into three steps (step 1, 2 and 3), and the general methods for synthesizing the respective compounds are as follows.

Step 1: General method for synthesizing compound (1-2)

Compound 1-1 (5.3 mmol) was dissolved in pyridine (10 mL), cooled to 0°C, and then acyl chloride (6.34 mmol) was slowly added thereto. The resulting reaction mixture was stirred at room temperature for 24 hours. The mixture was neutralized with aqueous 1M hydrochloric acid, diluted with water, and then extracted several times

with dichloromethane. The extracted organic layers were collected, washed successively with water and saturated aqueous sodium chloride solution, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure. The resulting residue was purified by column chromatography (eluent: hexane/ethyl acetate 5 = 1/1) to yield the title compound 1-2.

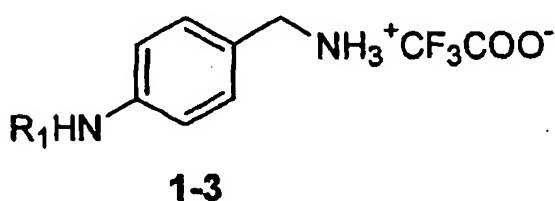


Example-step	Compound No.	R ₁	Spectral data
1-1	1-2a	SO ₂ CH ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.1-7.3 (m, 4H), 6.77 (s, 1H), 4.88 (bs, 1H), 4.28 (d, 2H), 2.99 (s, 3H), 1.46 (s, 9H)
2-1	1-2b	SO ₂ Et	¹ H NMR(300MHz, CDCl ₃) : δ 7.1-7.3 (m, 4H), 6.80 (s, 1H), 4.88 (bs, 1H), 4.25 (d, 2H), 3.07 (q, 3H), 1.46 (s, 9H), 1.36 (t, 3H)
3-1	1-2c	SO ₂ Pr	¹ H NMR(300MHz, CDCl ₃) : δ 7.23 (dd, 4H), 6.85 (s, 1H), 4.89 (bs, 1H), 4.28 (d, 2H), 3.05 (m, 2H), 1.67 (m, 2H), 1.46 (s, 9H), 1.01 (t, 3H)
4-1	1-2d	SO ₂ iPr	¹ H NMR(300MHz, CDCl ₃) : δ 7.17-7.31 (m, 4H), 6.98 (s, 1H), 4.87 (bs, 1H), 4.28 (d, 2H), 3.28 (m, 1H), 1.3-1.5 (m, 15H)
5-1	1-2e	SO ₂ Ph	¹ H NMR(300MHz, CDCl ₃) : δ 7.75 (d, 2H), 7.54 (m, 3H), 7.15 (d, 2H), 7.02 (d, 2H), 6.68 (s, 1H), 4.80 (t, 1H), 4.23 (d, 2H), 1.44 (s, 9H)

6-1	1-2f	SO ₂ CF ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.2-7.4 (m, 4H), 6.68 (s, 1H), 4.21 (d, 2H), 1.37 (s, 9H)
7-1	1-2g	COCH ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.46 (d, 2H), 7.24 (d, 2H), 4.80 (bs, 1H), 4.27 (d, 2H), 2.17 (s, 3H), 1.45 (s, 9H)
8-1	1-2h	COiPr	¹ H NMR(300MHz, CDCl ₃) : δ 7.50 (d, 2H), 7.24 (d, 2H), 7.11 (bs, 1H), 4.78 (bs, 1H), 4.25 (d, 2H), 2.50 (m, 1H), 1.45 (s, 9H), 1.24 (d, 6H)

Step 2: General method for synthesizing compound (1-3)

Compound **1-2** (4.8 mmol) was dissolved in dichloromethane (20 mL), cooled to 0°C, and trifluoroacetic acid (5 mL) was slowly added thereto. The resulting mixture was stirred at 0°C for an hour and 30 minutes. The solvent was evaporated under reduced pressure to obtain a flame-colored residue. The obtained residue was washed with ethyl ether and filtered to yield the title compound **1-3**.



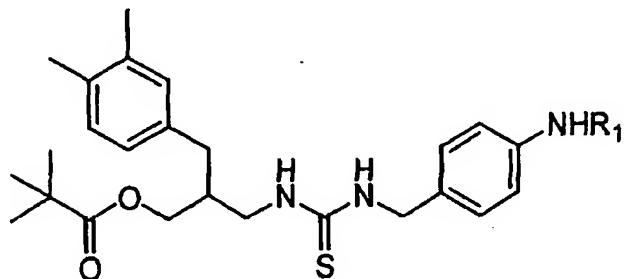
Example-step	Compound No.	R ₁	Spectral data

1-2	1-3a	SO ₂ CH ₃	¹ H NMR(300MHz, DMSO) : δ 8.14 (bs, 3H), 7.39 (d, 2H), 7.22 (d, 2H), 3.97 (s, 2H), 2.99 (s, 3H)
2-2	1-3b	SO ₂ Et	¹ H NMR(300MHz, DMSO) : δ 8.09 (bs, 3H), 7.51 (d, 2H), 7.33 (d, 2H), 4.01 (s, 2H), 2.79 (q, 2H), 1.46 (t, 3H)
3-2	1-3c	SO ₂ Pr	¹ H NMR(300MHz, DMSO) : δ 7.69 (d, 2H), 7.35 (d, 2H), 4.03 (s, 2H) 2.76 (t, 2H), 1.92 (m, 2H), 1.05 (t, 3H)
4-2	1-3d	SO ₂ iPr	¹ H NMR(300MHz, DMSO) : δ 7.82 (d, 2H), 7.37 (d, 2H), 3.99 (s, 2H) 3.53 (m, 1H), 1.53 (d, 6H)
5-2	1-3e	SO ₂ Ph	¹ H NMR(300MHz, DMSO) : δ 8.17 (d, 2H), 7.89 (d, 2H), 7.4-7.6 (m, 3H), 4.03 (s, 2H)
6-2	1-3f	SO ₂ CF ₃	¹ H NMR(300MHz, DMSO) : δ 7.9 (bs, 3H), 7.1-7.3 (m, 4H), 4.00 (bs, 2H)
7-2	1-3g	COCH ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.49 (m, 4H), 4.17 (s, 2H), 2.18 (s, 3H)
8-2	1-3h	COiPr	¹ H NMR(300MHz, CDCl ₃) : δ 7.50 (m, 4H), 4.17 (s, 2H), 2.70 (m, 1H), 1.22 (d, 6H)

Step 3: General method for synthesizing compound (1-4)

The compound **1-3** (0.38 mmol) was dissolved in DMF (1 mL), and triethylamine (0.38 mmol) was added thereto. The mixture was stirred at room temperature under nitrogen for 1 hour. To the mixture was added 2,2-dimethyl-propionic acid 3-(3,4-dimethylphenyl)-2-isothiocyanatomethylpropyl ester (0.38 mmol), and the resulting mixture was stirred for 24 hours. The reaction mixture was diluted with water, extracted several times with ethyl acetate. The collected organic layer was washed successively with water and saturated aqueous sodium chloride solution, dried over anhydrous magnesium sulfate, and concentrated under reduced

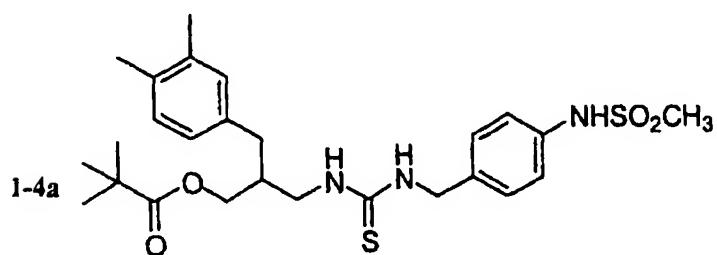
pressure to obtain a residue. The obtained residue was purified by column chromatography (eluent: hexane/ethyl acetate = 3/2) to yield the compound 1-4.

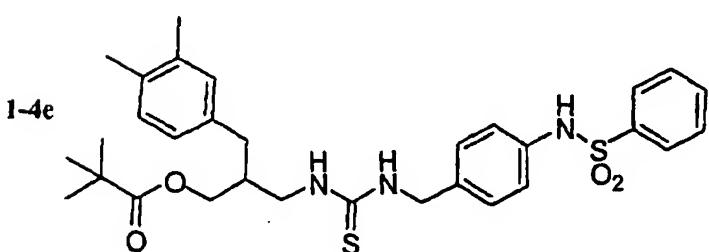
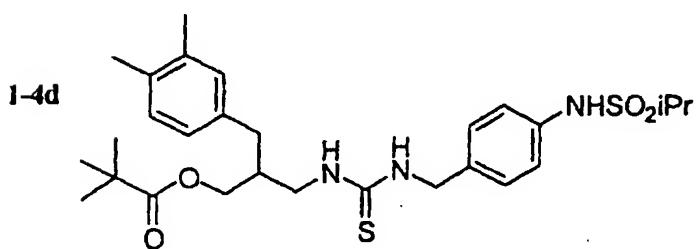
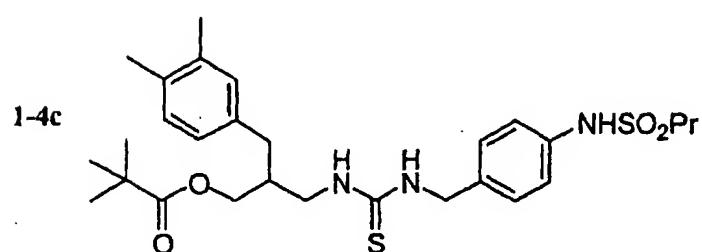
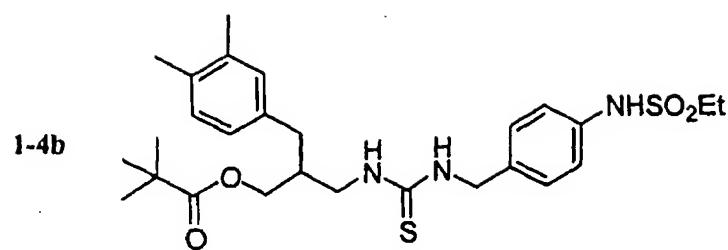


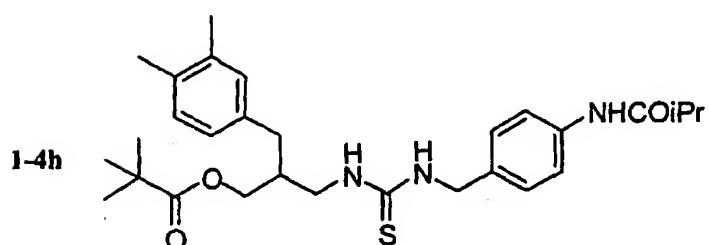
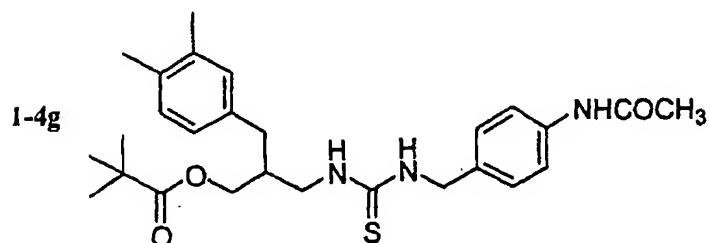
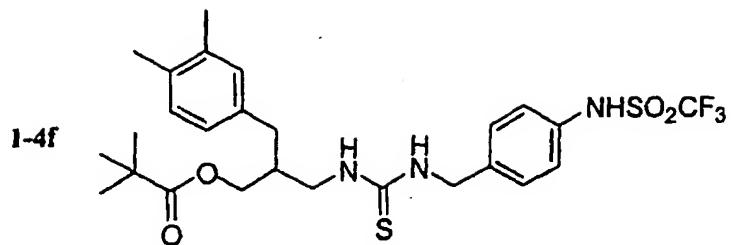
1-4

Example -step	Compound No.	R ₁	Spectral data
1-3	1-4a	SO ₂ CH ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.28 (d, 2H), 7.17 (d, 2H), 6.8-7.1 (m, 3H), 6.68 (s, 1H), 6.34 (m, 1H), 6.09 (brs, 1H), 4.51 (bs, 2H), 4.17 (m, 1H), 3.79 (m, 2H), 3.22 (m, 1H), 2.99 (s, 3H), 2.5-2.7 (m, 2H), 2.2-2.3 (m, 7H), 1.23 (s, 9H) ; mp=60 °C
2-3	1-4b	SO ₂ Et	¹ H NMR(300MHz, CDCl ₃) : δ 7.25 (d, 2H), 7.16 (d, 2H), 6.8-7.1 (m, 3H), 6.40 (m, 1H), 6.21 (bs, 1H), 4.50 (brs, 2H), 4.15 (m, 1H), 3.78 (m, 2H), 3.24 (m, 1H), 3.08 (q, 3H), 2.5-2.7 (m, 2H), 2.2-2.3 (m, 7H), 1.24 (s, 9H); mp=48 °C
3-3	1-4c	SO ₂ Pr	¹ H NMR(300MHz, CDCl ₃) : δ 7.25 (d, 2H), 7.14 (d, 2H), 6.8-7.1 (m, 3H), 6.45 (bs, 1H), 6.30 (brs, 1H), 4.51 (brs, 2H), 4.15 (m, 1H), 3.81 (m, 2H), 3.24 (m, 1H), 3.03 (t, 2H), 2.5-2.7 (m, 2H), 2.23 (m, 7H), 1.83 (q, 2H), 1.24 (s, 9H), 1.00 (t, 3H); mp=54 °C

4-3	1-4d	SO ₂ iPr	¹ H NMR(300MHz, CDCl ₃) : δ 7.1-7.3 (m, 4H), 6.8-7.0 (m, 3H), 6.48 (m, 2H), 4.50 (brs, 2H), 4.13 (m, 1H), 3.80 (m, 2H), 3.25 (m, 2H), 2.5-2.7 (m, 2H), 2.25 (m, 7H), 1.34 (d, 6H), 1.23 (s, 9H); mp=48 °C
5-3	1-4e	SO ₂ Ph	¹ H NMR(300MHz, CDCl ₃) : δ 7.76 (d, 2H), 7.52 (m, 2H), 6.8-7.1 (m, 3H), 6.41 (bs, 2H), 6.19 (brs, 2H), 4.43 (brs, 2H), 4.12 (m, 1H), 3.78 (m, 2H), 3.21 (m, 1H), 2.5-2.7 (m, 2H), 2.23 (m, 7H), 1.23 (s, 9H); mp=66 °C
6-3	1-4f	SO ₂ CF ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.28 (d, 2H), 7.17 (d, 2H), 6.8-7.1 (m, 3H), 6.48 (brs, 1H), 6.36 (brs, 1H), 4.50 (brs, 2H), 4.15 (m, 1H), 3.75 (m, 2H), 3.28 (m, 1H), 2.58 (m, 2H), 2.11-2.24 (m, 7H), 1.23 (s, 9H); mp=44 °C
7-3	1-4g	COCH ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.46 (d, 2H), 7.24 (d, 2H), 6.9 (m, 3H), 6.28 (brs, 1H), 6.07 (brs, 1H), 4.43 (bs, 2H), 4.13 (m, 1H), 3.75 (m, 2H), 3.27 (m, 1H), 2.54 (m, 2H), 2.21 (m, 10H), 1.22 (s, 9H)
8-3	1-4h	COiPr	¹ H NMR(300MHz, CDCl ₃) : δ 7.51 (d, 2H), 7.22 (m, 2H), 6.93 (m, 3H), 6.28 (brs, 1H), 6.07 (brs, 1H), 4.41 (brs, 2H), 4.13 (m, 1H), 3.74 (m, 2H), 3.27 (m, 1H), 2.52 (m, 3H), 2.21 (m, 7H), 1.24 (m, 15H)







5 Examples 9~17

Step 1: Synthesis of 4-chloromethyl-2-methoxy-1-nitrobenzene (2-1)

(3-methoxy-4-nitrophenyl)methanol (4.43 mmol) was dissolved in dichloromethane (10 mL), and then triethylamine (13.3 mmol) was added thereto. After the reaction mixture was cooled to 0 °C, methanesulfonyl chloride (6.64 mmol)

was slowly added thereto. The resulting mixture was stirred at room temperature under nitrogen for 24 hours. To the mixture was added ammonium chloride solution to terminate the reaction, diluted with water, and then extracted several times with dichloromethane. The collected organic layer was washed with water and brine, dried 5 over anhydrous magnesium sulfate, and concentrated under reduced pressure. The resulting residue thus obtained was purified by column chromatography (eluent: hexane/ethyl acetate = 4/1) to yield the title compound **2-1** (yield: 93%) as a yellow solid.

¹H NMR (CDCl₃) δ 7.86 (d, 1H), 7.05 (s, 1H), 6.98 (d, 1H), 4.56 (s, 2H), 3.99 (s, 3H)

10

Step 2: Synthesis of 4-azidomethyl-2-methoxy-1-nitro-benzene (**2-2**)

A solution of the compound **2-1** (4.12 mmol) and sodium azide (16.5 mmol) in dimethylformamide (3 mL) was stirred at room temperature under nitrogen for 1 hour. The reaction mixture was diluted with water, extracted several times with ethyl acetate, 15 and then the organic layer was collected. The collected organic layer was washed successively with water and saturated aqueous sodium chloride solution, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. The resulting residue was purified by column chromatography (eluent: hexane/ethyl acetate = 3/1) to

yield the title compound **2-2** (yield: 95%) as a yellow solid.

¹H NMR (CDCl₃) δ 7.87 (d, 1H), 7.05 (s, 1H), 6.99 (d, 1H), 4.45 (s, 2H), 3.99 (s, 3H)

Step 3: Synthesis of (3-methoxy-4-nitro-benzyl)-carbamic acid t-butyl ester (**2-3**)

5 A solution of the compound **2-2** (3.91 mmol), triphenyl phosphine (7.83 mmol) and water (7.83 mmol) in THF (30 mL) was stirred at room temperature for 24 hours. The reaction mixture was concentrated under reduced pressure. The residue thus obtained was dissolved in ethanol (20 mL), and Boc₂O (1.71 g) was added thereto. The resulting mixture was stirred at room temperature for 2 hours. After the reaction 10 mixture was concentrated under reduced pressure, the resulting residue was purified by column chromatography (eluent: hexane/ethyl acetate = 2/1) to yield the title compound **2-3** (yield: 86%) as a yellow liquid.

¹H NMR (CDCl₃) δ 7.83 (d, 1H), 7.02 (s, 1H), 6.93 (d, 1H), 5.01 (brs, 1H), 4.35 (d, 2H), 3.95 (s, 3H), 1.47 (s, 9H)

15

Step 4: Synthesis of (4-amino-3-methoxybenzyl)carbamic acid t-butyl ester (**2-4**)

The compound **2-3** (3.13 mmol) was dissolved in methanol (20 mL), and then

10% palladium/carbon catalyst (85 mg) was added thereto. The resulting mixture was stirred at room temperature under hydrogen for 1 hour. After the reaction mixture was filtered, the filtrate was concentrated under reduced pressure to yield the title compound **2-4** (yield: 90%) as a brown liquid.

5 ^1H NMR (CDCl₃) δ 6.6-6.7 (m, 3H), 4.19 (d, 2H), 3.88 (s, 3H), 1.46 (s, 9H)

Step 5: Synthesis of (4-methanesulfonylamino-3-methoxybenzyl)carbamic acid t-butyl ester (**2-5**)

The title compound **2-5** was obtained as a pink solid in a yield of 67%
10 according to the same procedure as synthesizing method of compound 1-2a, except that the compound **2-4** (3.13 mmol) was used as a starting material.

^1H NMR (CDCl₃) δ 7.47 (d, 1H), 6.86 (m, 2H), 6.74 (s, 1H), 4.88 (bs, 1H), 3.88 (s, 3H), 2.94 (s, 3H), 1.47 (s, 9H)

15 Step 6: Synthesis of 4-methanesulfonylamino-3-methoxybenzylammonium trifluoroacetate (**2-7**)

The title compound **2-7** was obtained as a green solid in a yield of 87%

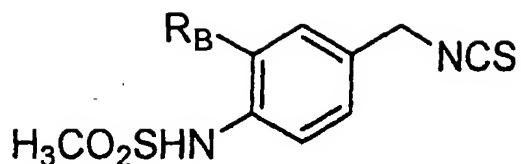
according to the same procedure as synthesizing method of compound **1-3a**, except that the compound **2-5** (2.1 mmol) was used as a starting material.

¹H NMR (DMSO) δ 8.16 (bs, 3H), 7.29 (d, 1H), 7.20 (s, 1H), 6.99 (d, 1H), 4.00 (s, 2H), 3.82 (s, 3H), 2.95 (s, 3H)

5

Step 7: General method for synthesizing compounds (2-8 and 2-9)

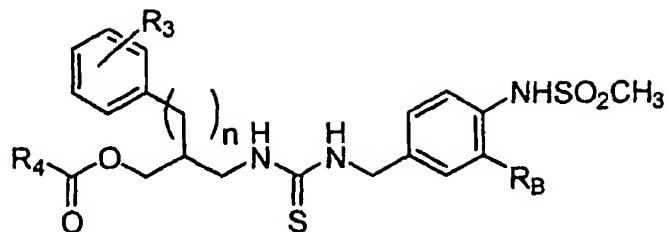
The compound **2-6** or **2-7** (12 mmol) was dissolved in DMF (1mL), and then triethylamine (12 mmol) was added thereto. The resulting mixture was stirred at room temperature under nitrogen for 1 hour. To the mixture was added 1,1-thio-1H-2-carbonyl-di-2-pyridone (12 mmol), and stirred for 24 hours. The reaction mixture was diluted with water, extracted several times with ethyl acetate, and then the organic layer was collected. The collected organic layer was washed with water and then with saturated aqueous sodium chloride solution, dried over anhydrous magnesium sulfate, and concentrated under reduced pressure. The resulting residue was purified by column chromatography (eluent: hexane/ethyl acetate = 3/2) to yield the title compounds **2-8** or **2-9**.

**2-8, 2-9**

Compound No.	R _B	Spectral data
2-8	H	¹ H NMR(300MHz, CDCl ₃) : δ 7.32 (d, 2H) 7.25 (d, 2H), 6.62 (s, 1H), 4.70 (s, 2H) 3.04 (s, 3H)
2-9	OCH ₃	¹ H NMR(300MHz, CDCl ₃) : δ 7.29 (s, 1H) 6.96 (m, 2H), 5.09 (s, 2H), 3.82 (s, 3H), 3.04 (s, 3H)

Step 8: General method for synthesizing compounds (2-10 ~ 2-18)

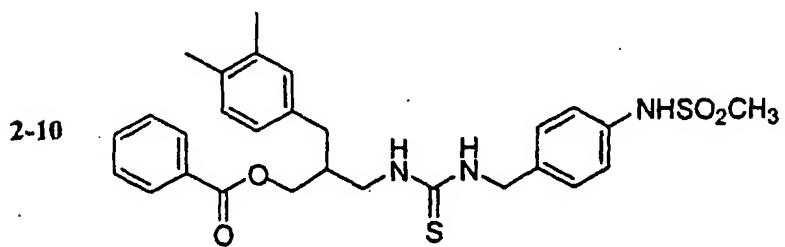
5 A solution of azide compound (0.5 mmol) prepared in accordance with the well-known method, Lindlers catalyst (50 mmol) and the compound 2-8 in ethanol (5 mL) was subjected to hydrogenation reaction under hydrogen atmosphere for 2 hours. After the reaction mixture was filtered, the filtrate was concentrated under reduced pressure. The concentrate was purified by column chromatography (eluent: ethyl acetate/hexane = 1/1) to yield the title compounds 2-10 ~ 2-18

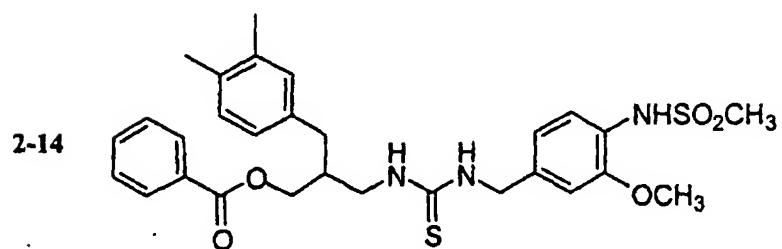
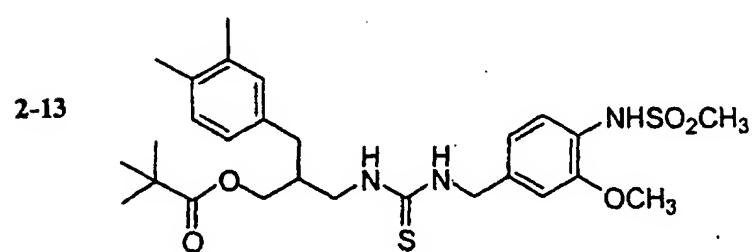
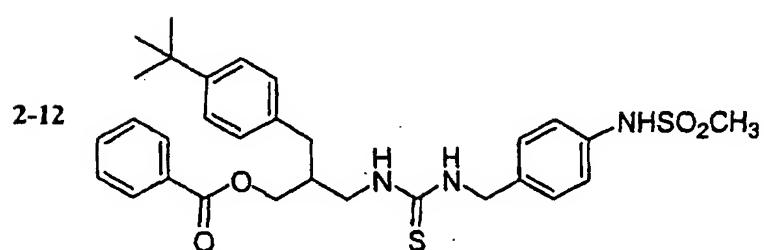
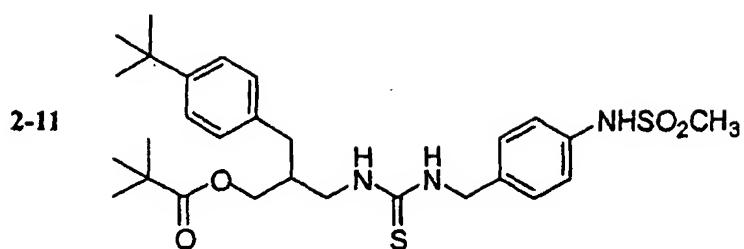


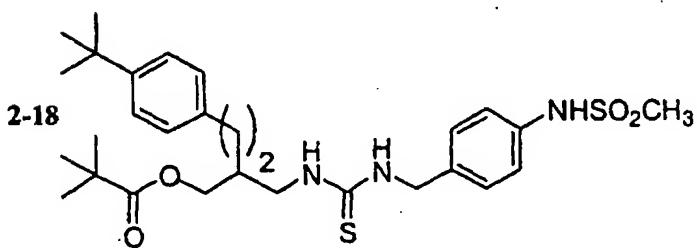
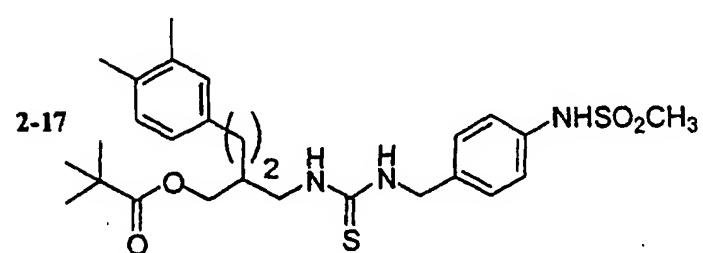
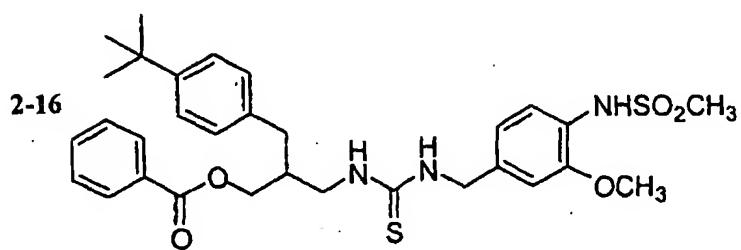
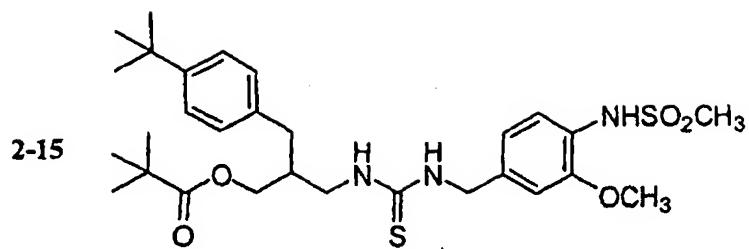
2-10 ~ 2-18

Exam ples	Compound No.	R	Spectral data
9	2-10	$R_B = H$ $R_3 = 3,4\text{-di-Me}$ $R_4 = Ph$ $n=1$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 8.01 (m, 2H), 7.4-7.6 (m, 3H), 7.1-7.3 (m, 4H), 6.9-7.0 (m, 3H), 6.53 (m, 1H), 6.40 (brs, 1H), 4.52 (brs, 2H), 4.35 (m, 1H), 4.17 (m, 1H), 3.82 (m, 1H), 3.41 (m, 1H), 2.93 (s, 3H), 2.5-2.7 (m, 2H), 2.1-2.3 (m, 7H); mp=46°C
10	2-11	$R_B = H$ $R_3 = 4\text{-t-Bu}$ $R_4 = t\text{-Bu}$ $n=1$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 7.31 (d, 4H), 7.18 (d, 2H), 7.10 (d, 2H), 6.51 (s, 1H), 6.33 (t, 1H), 6.03 (brs, 1H), 4.53 (brs, 2H), 4.15 (m, 1H), 3.82 (m, 2H), 3.23 (m, 1H), 2.99 (s, 3H), 2.60 (t, 2H), 2.32 (m, 1H), 1.25 (d, 18H); mp=61°C
11	2-12	$R_B = H$ $R_3 = 4\text{-t-Bu}$ $R_4 = Ph$ $n=1$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 8.01 (d, 2H), 7.4-7.6 (m, 3H), 7.1-7.3 (m, 4H), 6.54 (s, 1H), 6.39 (t, 1H), 6.10 (brs, 1H), 4.53 (brs, 2H), 4.39 (m, 1H), 4.07 (m, 1H), 3.82 (m, 1H), 3.38 (m, 1H), 2.97 (s, 3H), 2.68 (t, 2H), 2.48 (m, 1H, CH), 1.29 (s, 9H); mp=68°C
12	2-13	$R_B = OMe$ $R_3 = 3,4\text{-di-Me}$ $R_4 = t\text{-Bu}$ $n=1$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 7.48 (d, 2H), 6.9-7.1 (m, 5H), 6.77 (s, 1H), 6.31 (m, 1H), 5.99 (brs, 1H), 4.49 (bs, 2H), 4.20 (m, 1H), 3.88 (s, 3H), 3.85 (m, 2H), 3.22 (m, 1H), 2.93 (s, 3H), 2.56 (m, 2H), 2.2-2.3 (m, 7H), 1.23 (s, 9H); mp=49°C
13	2-14	$R_B = OMe$ $R_3 = 3,4\text{-di-Me}$ $R_4 = Ph$ $n=1$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 8.02 (t, 2H), 7.4-7.6 (m, 3H), 6.9-7.1 (m, 3H), 6.77 (s, 1H), 6.38 (m, 1H), 6.09 (bs, 1H), 4.45 (m, 3H), 4.05 (m, 1H), 3.85 (s, 3H), 3.37 (m, 1H), 2.92 (s, 3H), 2.67 (m, 2H), 2.43 (m, 1H), 2.25 (m, 6H); mp=62°C

14	2-15	$R_B = OMe$ $R_3 = 4-t\text{-Bu}$ $R_4 = t\text{-Bu}$ $n=1$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 7.45 (d, 2H), 7.30 (d, 2H), 7.08 (d, 2H), 6.8-7.0 (m, 3H), 6.39 (t, 1H), 6.24 (bs, 1H), 4.52 (bs, 2H), 4.15 (m, 1H), 3.82 (m, 5H), 3.25 (m, 1H) 2.93 (s, 3H), 2.60 (m, 2H), 2.31 (m, 1H), 1.25 (d, 18H); mp=62°C
15	2-16	$R_B = OMe$ $R_3 = 4-t\text{-Bu}$ $R_4 = Ph$ $n=1$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 8.03 (d, 2H), 7.4-7.6 (m, 3H), 7.1-7.3 (m, 4H), 6.77 (s, 1H), 6.38 (t, 1H), 6.10 (bs, 1H), 4.51 (bs, 2H), 4.43 (m, 1H), 4.08 (m, 1H), 3.82 (m, 4H), 3.38 (m, 1H), 2.92 (s, 3H), 2.69 (t, 2H), 2.36 (m, 1H), 1.29 (s, 9H); mp=57°C
16	2-17	$R_B = H$ $R_3 = 3,4\text{-di-Me}$ $R_4 = t\text{-Bu}$ $n=2$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 7.1-7.3 (m, 7H), 6.39 (t, 1H), 6.06 (bs, 1H), 4.59 (d, 2H), 4.16 (m, 1H), 3.97 (m, 1H) 3.77 (m, 2H), 3.22 (m, 1H), 2.99 (s, 3H), 2.68 (t, 2H), 2.05 (m, 1H), 1.58 (m, 2H), 1.25 (d, 9H)
17	2-18	$R_B = H$ $R_3 = 4-t\text{-Bu}$ $R_4 = t\text{-Bu}$ $n=2$	$^1\text{H NMR}$ (300MHz, CDCl_3) : δ 7.33 (d, 2H), 7.17 (d, 2H), 6.9-7.1 (m, 3H), 6.42 (s, 1H), 6.36 (t, 1H), 6.06 (bs, 1H), 4.59 (d, 2H), 4.21 (m, 1H), 3.97 (m, 1H) 3.72 (m, 1H), 3.22 (m, 2H), 2.99 (s, 3H), 2.65 (t, 2H), 2.23 (m, 7H), 1.60 (m, 2H), 1.23 (s, 9H)

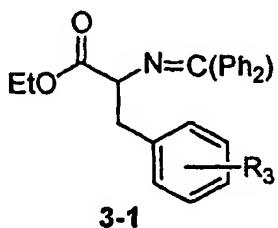






Step 1: General method for synthesizing compound (3-1)

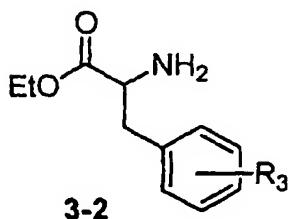
N-(diphenylmethyleneglycine)ethyl ester (1.83 mmol) was dissolved in dichloromethane (5mL), and then 50% aqueous sodium hydroxide solution (3.66 mmol), tetrabutylammonium bromide (1.83 mmol) and alkyl halide (2.50 mmol) were added thereto. The reaction mixture was stirred at room temperature for 24 hours, neutralized with 6N aqueous hydrochloric acid, diluted with water, and then extracted several times with dichloromethane. After the obtained organic layer was dried over anhydrous magnesium sulfate and filtered, the filtrate was concentrated under reduced pressure. The concentrate was purified by column-chromatography (eluent: hexane/ethyl acetate = 10/1) to yield the title compound 3-1.



Example-step	Compound No.	R ₃	Spectral data
18-1	3-1a	3,4-di-Me	¹ H NMR(300MHz, CDCl ₃) : δ 7.59 (t, 2H), 7.2-7.4 (m, 10H), 6.95 (m, 1H), 4.16-4.28 (m, 3H), 3.35 (dd, 1H), 3.17 (m, 1H), 2.12 (m, 6H), 1.23 (q, 3H)
19-1	3-1b	4-t-Bu	¹ H NMR(300MHz, CDCl ₃) : δ 7.59 (d, 2H), 7.2-7.4 (m, 10H), 6.95 (d, 2H), 4.11-4.24 (m, 3H), 3.1-3.3 (m, 2H), 1.25 (m, 12H)

Step 2: General method for synthesizing compound (3-2)

After the compound 3-1 (1.18 mmol) was dissolved in THF (10 ml), the
 5 solution was adjusted to a pH 4 with 1N aqueous hydrochloric acid. The mixture was
 stirred at room temperature for 30 minutes. To the mixture was added ethyl acetate
 (10 mL) and water (10 mL). The formed aqueous layer was neutralized to a pH 9 with
 1N aqueous sodium hydroxide solution, and then extracted several times with ethyl
 acetate. After the organic layer thus obtained was dried over anhydrous magnesium
 10 sulfate and filtered, the filtrate was concentrated under reduced pressure. The
 concentrate was purified by column chromatography (eluent: hexane/ethyl acetate =
 2/1) to yield the title compound 3-2.

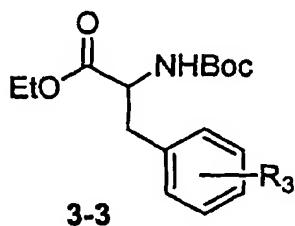


Example- step	Compound No.	R ₃	Spectral data
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18-2	3-2a	3,4-di-Me	¹ H NMR(300MHz, CDCl ₃) : δ 6.9-7.1 (m, 3H), 4.19 (q, 2H), 3.73 (m, 1H), 3.55 (dd, 1H), 3.17 (dd, 1H), 2.22 (m, 6H), 1.24 (m, 3H)
19-2	3-2b	4-t-Bu	¹ H NMR(300MHz, CDCl ₃) : δ 7.33 (d, 2H), 7.24 (d, 2H), 4.18 (q, 2H), 3.70 (m, 1H), 3.05 (dd, 1H), 2.83 (dd, 1H), 1.25 (m, 12H)

Step 3: General method for synthesizing compound (3-3)

After the compound 3-2 (0.68 mmol) was dissolved in dichloromethane (3 mL),
 5 the solution was cooled to 0 °C and (Boc)₂O (0.75 mmol) was added thereto. The reaction mixture was stirred at room temperature for 24 hours. The mixture was concentrated under reduced pressure. The resulting residue was purified by column chromatography (eluent: hexane/ethyl acetate = 4/1) to yield the title compound 3-3.

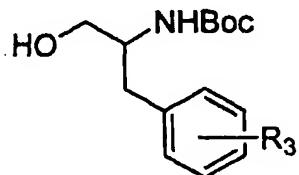


Example-step	Compound No.	R ₃	Spectral data
18-3	3-3a	3,4-di-Me	¹ H NMR(300MHz, CDCl ₃) : δ 6.9-7.2 (m, 3H), 4.97 (m, 1H), 4.50 (m, 1H), 4.15 (q, 2H), 3.05 (m, 2H), 2.22 (m, 6H), 1.42 (s, 9H), 1.20 (t, 3H)

19-3	3-3b	4-t-Bu	¹ H NMR(300MHz, CDCl ₃) : δ 7.31 (d, 2H), 7.06 (d, 2H), 4.56 (m, 1H), 4.18 (q, 2H), 3.05 (d, 2H), 1.41 (s, 9H), 1.29 (s, 9H), 1.19 (t, 3H)
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Step 4: General method for synthesizing compound (3-4)

After lithium aluminum hydride (1.79 mmol) was suspended in diethyl ether (5 mL), the suspension was cooled to 0°C and a solution of the compound 3-3 (0.48 mmol) in diethyl ether (5 mL) was added thereto by dropping. After the reaction mixture was stirred at room temperature for 5 hours, the mixture was cooled to 0°C again. Water (700 μl), NaOH (1.4 mL) and water (2.1 mL) were added to the mixture in succession, and stirred continuously. After the resulting mixture was washed with ethyl acetate and filtered, the obtained filtrate was concentrated under reduced pressure. The resulting residue was purified by column chromatography (eluent: ethyl acetate/hexane = 4/1) to yield the title compound 3-4.

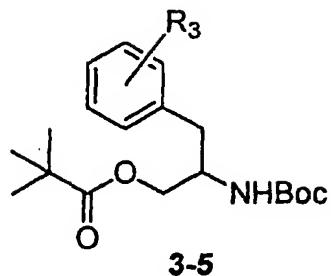


3-4

Example-step	Compound No.	R ₃	Spectral data
18-4	3-4a	3,4-di-Me	¹ H NMR(300MHz, CDCl ₃) : δ 6.9-7.2 (m, 3H), 4.71 (bs, 1H), 3.83 (m, 1H), 3.5-3.7 (m, 2H), 2.85 (dd, 2H), 2.24 (m, 6H), 1.41 (s, 9H)
19-4	3-4b	4-t-Bu	¹ H NMR(300MHz, CDCl ₃) : δ 7.32 (d, 2H), 7.13 (d, 2H), 4.71 (bs, 1H), 3.87 (m, 1H), 3.5-3.7 (m, 2H), 2.79 (d, 2H), 1.41 (s, 9H), 1.30 (s, 9H)

Step 5: General method for synthesizing compound (3-5)

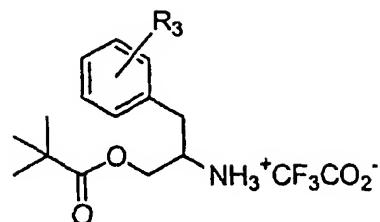
A mixture of the compound 3-4 (0.71 mmol), triethylamine (2.8 mmol) and a catalytic amount of 4-dimethylaminopyridine (DMAP) was dissolved in dichloromethane (7 mL), cooled to 0°C, and then trimethylacetyl chloride (2.8 mmol) was added thereto. The reaction mixture was stirred at room temperature for 1 hour. The mixture was diluted with dichloromethane, washed with 1N HCl, water and saturated aqueous sodium chloride solution once in succession, dried over anhydrous magnesium sulfate and filtered. The obtained filtrate was then concentrated under reduced pressure. The resulting residue was purified by column chromatography (eluent: ethyl acetate/hexane = 1/4) to yield the title compound 3-5.



Example-step	Compound No.	R_3	Spectral data
18-5	3-5a	3,4-di-Me	^1H NMR(300MHz, CDCl_3) : δ 6.9-7.2 (m, 3H), 4.61 (bs, 1H), 4.01 (m, 3H), 2.86 (m, 2H), 2.24 (m, 6H), 1.41 (m, 9H), 1.23 (m, 9H)
19-5	3-5b	4-t-Bu	^1H NMR(300MHz, CDCl_3) : δ 7.30 (d, 2H), 7.10 (d, 2H), 4.57 (bs, 1H), 4.13 (m, 3H), 2.78 (m, 2H), 1.41 (m, 9H), 1.2-1.3 (m, 18H)

Step 6: General method for synthesizing compound (3-6)

5 The compound **3-5** (0.71 mmol) was dissolved in dichloromethane (6 mL), cooled to 0°C, and then trifluoroacetic acid (1.5 mL) was slowly added thereto. The reaction mixture was stirred at room temperature for 3 hours. The mixture was concentrated under reduced pressure to yield the unpurified title amine salt **3-6**.



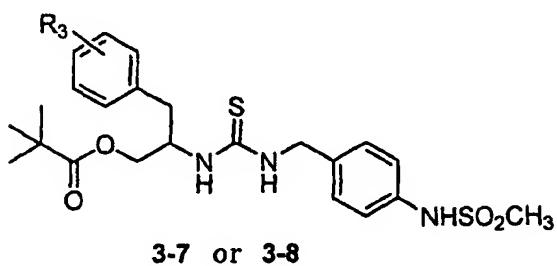
3-6

Example-step	Compound No.	R ₃	Spectral data
18-6	3-6a	3,4-di-Me	¹ H NMR(300MHz, CDCl ₃) : δ 7.37 (d, 1H), 7.13 (m, 2H), 4.1-4.3 (m, 2H), 3.71 (m, 1H), 3.04 (m, 2H), 2.22 (m, 6H), 1.22 (m, 9H)
19-6	3-6b	4-t-Bu	¹ H NMR(300MHz, CDCl ₃) : δ 7.35 (m, 2H), 7.13 (m, 2H), 4.1-4.2 (m, 3H), 2.80 (d, 2H), 1.30 (s, 9H), 1.09 (s, 9H)

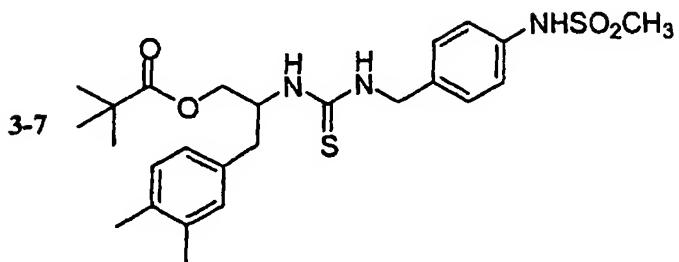
Step 7: Method for synthesizing compounds (3-7 and 3-8)

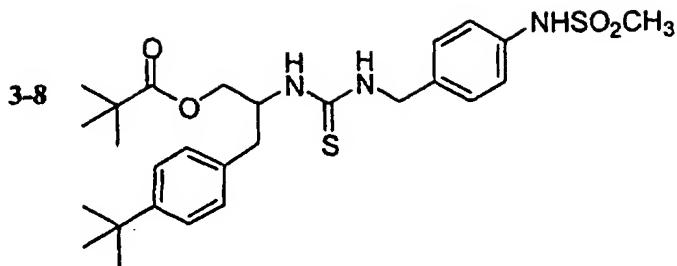
5 To a solution of compound 3-6 (0.46 mmol) in dimethylformamide (1 mL) was added triethylamine (0.46 mmol). The resulting mixture was stirred at room temperature under nitrogen for 30 minutes. To the mixture was added the compound 2-8 (0.46 mmol), and stirred at room temperature for 24 hours. The reaction mixture was diluted with water, and extracted several times with ethyl acetate. The obtained 10 organic layer was washed with water and brine, dried over anhydrous magnesium sulfate, and then concentrated under reduced pressure. The resulting residue was

purified by column chromatography (eluent: hexane/ethyl acetate = 1/1) to yield the title compounds **3-7 or 3-8**.



Example step	Compound No.	R ₃	Spectral data
18-7	3-7	3,4-di-Me	¹ H NMR(300MHz, CDCl ₃) : δ 7.19 (m, 4H), 7.04 (m, 3H), 6.28 (brs, 1H), 6.07 (brs, 1H), 4.56 (m, 2H), 4.17 (m, 1H), 3.99 (m, 1H), 3.01 (m, 5H), 2.76 (m, 1H), 2.26 (m, 6H), 1.19 (s, 9H)
19-7	3-8	4-t-Bu	¹ H NMR(300MHz, CDCl ₃) : δ 7.27 (m, 8H), 6.68 (brs, 1H), 6.51 (brs, 1H), 6.11 (m, 1H), 4.58 (m, 2H), 4.20 (m, 1H), 3.99 (m, 1H), 3.12 (m, 5H), 2.76 (m, 1H), 1.30 (m, 18H)

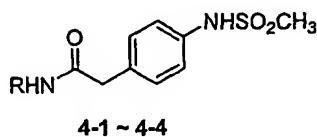




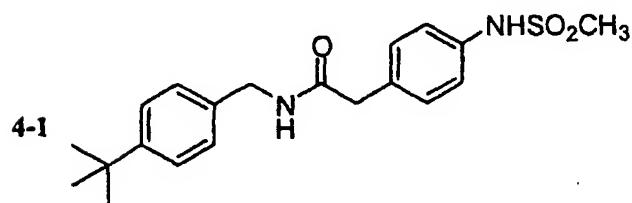
Examples 20~23

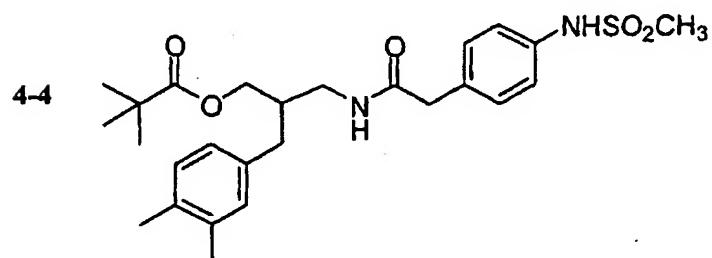
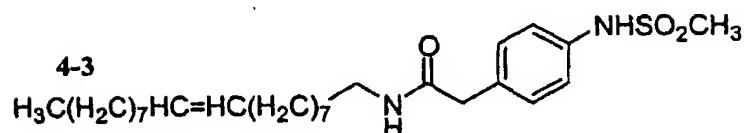
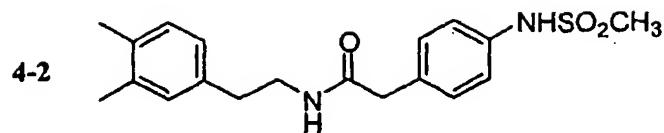
General method for synthesizing compounds (4-1 ~ 4-4)

5 A solution of amine compound (0.5 mmol) and
 (4-methanesulfonylaminophenyl)acetic acid pentafluorophenyl ester (0.5 mmol) in
 dichloromethane was stirred at room temperature for 24 hours. The reaction mixture
 was concentrated under reduced pressure. The resulting residue was purified by
 column chromatography (eluent: hexane/ethyl acetate = 1/2) to yield the title
 10 compounds 4-1 ~ 4-4.



Examples	Compound No.	R	Spectral data
20	4-1	4-t-BuPhCH ₂ -	¹ H NMR(300MHz, CDCl ₃) : δ 7.1-7.3 (m, 8H), 6.44 (brs, 1H), 5.66 (brs, 1H), 4.40 (d, 2H), 3.57 (s, 2H), 3.00 (s, 3H), 1.30 (s, 9H);mp=54°C
21	4-2	3,4-di-Ph(CH ₂) ₂ -	¹ H NMR(300MHz, CDCl ₃) : δ 7.1-7.3 (m, 4H), 6.43 (brs, 1H), 5.35 (m, 3H), 3.52 (s, 2H), 3.21 (q, 2H), 3.02 (s, 3H), 2.00 (d, 4H) 1.56 (s, 2H), 1.26 (s, 22H), 0.88 (t, 3H);mp=134°C
22	4-3	Oleyl	¹ H NMR(300MHz, CDCl ₃) : δ 7.23 (m, 4H), 7.04 (d, 1H), 6.87 (m, 2H), 6.38 (brs, 1H), 5.32 (brs, 1H), 3.49 (s, 2H), 3.25 (q, 2H), 3.01 (s, 3H), 2.52 (t, 2H), 2.22 (s, 6H), 1.77 (t, 2H)
23	4-4		¹ H NMR(300MHz, CDCl ₃) : δ 7.1-7.3 (m, 4H), 6.8-7.1 (m, 3H), 5.90 (t, 1H), 4.13 (m, 1H), 3.80 (m, 1H), 3.54 (s, 2H), 3.35 (m, 1H), 3.11 (m, 1H), 2.97 (s, 3H), 2.5-2.7 (m, 2H), 2.1-2.3 (m, 7H), 1.24 (s, 9H);mp=30°C





In the below, it is confirmed by calcium influx test that the compounds of the present invention are antagonist to vanilloid receptor, further confirmed by analgesic effect test that they exhibit strong analgesic effects while showing no irritation shown in an agonist.

Experimental Example. Biological potency test

10 (1) ^{45}Ca influx test

1) Separation of spinal dorsal root ganglia (DRG) in newborn rats and primary

culture thereof

Neonatal(2-day old or younger than 2-day old) SD rats were put in ice for 5 minutes to anesthetize and disinfected with 70% ethanol. DRG of all part of spinal cord were dissected (Wood et al., 1988, J. Neurosci. 8, pp3208-3220) and collected in 5 DME/F12 medium to which 1.2 g/l sodium bicarbonate, 50 mg/l gentamycin were added. The DRG were incubated sequentially at 37°C for 30 min in 200 U/ml collagenase and 2.5 mg/ml trypsin, separately. The ganglia were washed twice with DME/F12 medium supplemented with 10% horse serum, triturated through a fire-polished Pasteur pipette, filtered through Nitex 40 membrane to obtain single cell 10 suspension. This was subjected to centrifugation, then re-suspended in cell culture medium at certain level of cell density. As the cell culture medium, DME/F12 medium supplemented with 10% horse serum, diluted 1:1 with identical medium conditioned by C6 glioma cells (2 days on a confluent monolayer) was used, and NGF(Nerve Growth Factor) was added to final concentration of 200 ng/ml. After the 15 cells were grown 2 days in medium where cytosine arabinoside (Ara-C, 100 µM) was added to kill dividing nonneuronal cells, medium was changed to one without Ara-C. The resuspended cells were plated at a density of 1500-1700 neurons/well onto Terasaki plates previously coated with 10 µg/ml poly-D-ornithine.

2) ^{45}Ca influx experiments

DRG nerve cells from the primary culture of 2-3 days were equilibrated by washing 4 times with HEPES (10mM, pH 7.4)-buffered Ca^{2+} , Mg^{2+} -free HBSS (H-HBSS). The solution in each well was removed from the individual well. Medium containing the test compound plus capsaicin (final concentration 0.5 μM) and ^{45}Ca (final concentration 10 $\mu\text{Ci/ml}$) in H-HBSS was added to each well and incubated at room temperature for 10 min. Terasaki plates were washed six times with H-HBSS and dried in an oven. To each well, 0.3% SDS (10 μl) was added to elute ^{45}Ca . After the addition of 2ml of scintillation cocktail into each well, the amount of ^{45}Ca influx into neuron was measured by counting radioactivity. Antagonistic activities of test compounds against vanilloid receptor were calculated as percent of the maximal response of capsaicin at a concentration of 0.5 μM and results are given as IC_{50} (Table 1).

(2) Channel activity assay

Antagonistic activities of test compounds were assayed based on electrical change of cation channel connected to vanilloid receptor and experiments were conducted according to reference method (Oh et al., 1996, J. Neuroscience 16, pp1659-1667) (Table 1).

Table 1. Results of Calcium Influx and Patchclamp Test

Examples	Calcium Uptake Test(IC ₅₀)	Patchclamp Test (antagonistic activities)
1	0.45	++
2	2.5	
3	4.3	
4	32	
5	>50	
6	11	
7	NR	
8	NR	
9	13	
10	2.5	
11	>50	
12	0.68	
13	38	
14	2.8	
15	>50	
16	1.1	
17	1.8	

NR: no response

+: antagonistic potency equal to capsazepine

++: antagonistic potency 10 times higher than capsazepine

(3) Analgesic activity test: Mouse writhing test by inducing with phenyl-p-quinone

Male ICR mice (mean body weight 25g) were maintained in a controlled
5 lighting environment (12 h on/ 12 h off) for experiment. Animals received an
intraperitoneal injection of 0.3ml of the chemical irritant phenyl-p-quinone (dissolved in
saline containing 5% ethanol to be a dose of 4.5mg/kg) and 6 min later, the number of
abdominal constrictions was counted in the subsequent 6 min period. Animals (10
animals/group) received 0.2ml of test compounds solution in vehicle of ethanol/Tween
10 80/saline (10/10/80) intraperitoneally 30 min before the injection of phenyl-p-quinone.
A reduction in the number of writhes responding to the test drug compound relative to
the number responding in saline control group was considered to be indicative of an
analgesic effect. Analgesic effect was calculated by % inhibition equation (%
15 inhibition=(C-T)/C x 100), wherein C and T represent the number of writhes in control
and compound-treated group, respectively (Table 2).

The test results demonstrated that analgesic effect of the compounds used in
this experiment is potent, and in particular, it is significant to clarify that vanilloid
receptor antagonist can exhibit such potent analgesic effect, and the results suggests that

vanilloid receptor antagonist has potential as an analgesic agent.

Table 2. Test result of analgesic activity for writhing by phenyl-p-quinone

Examples	Dose (mg/kg)	Analgesic effect (% Inhibition)
1	1	50
12	1	65

5 (4) Antiinflammatory activity test: TPA(12-O-tetradecanoylphorbol 13-acetate)-induced mouse ear edema test

Male ICR mice(body weight 25-30g), 10 animals/group, were treated topically on the right ear with 30 μ l of TPA (2.5 μ g) solution in acetone and after 15 min, 30 μ l of acetone or test compound solution in acetone was applied topically. After six hours, 10 an identical treatment was applied again. After twenty four hours following the treatment of TPA, the animals were sacrificed and ear tissue was dissected using 6 mm-diameter punch. Ear tissue dissected were weighed to the nearest 0.1 mg on an electrobalance. The increased weight of the tissue compared to control group was considered as an index of inflammation. The percent inhibition is defined by the 15 following equation:

% inhibition = $(C-T)/C \times 100$, wherein C and T represent an increase of ear weight in TPA-treated and TPA+drug-treated group, respectively.

The above experiment shows that vanilloid receptor antagonist exhibits significant anti-inflammatory effects. This phenomenon can be understood by connecting with the action of vanilloid receptor in neurogenic inflammation, and suggests potential applicability of vanilloid receptor antagonist in various inflammatory diseases, in particular, neurogenic inflammatory diseases.

Table 3. TPA-induced mice ear edema test

Examples	Dose (mg/ear)	Anti-inflammatory effect (% Inhibition)
1	1	34
12	1	41

10

Industrial Applicability

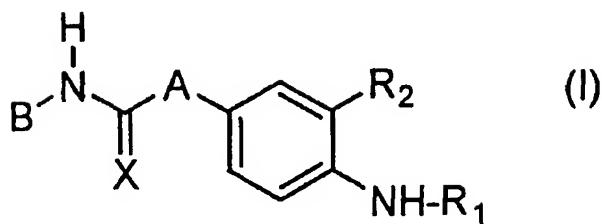
The compounds according to the present invention are useful in the prevention or treatment of pain, acute pain, chronic pain, neuropathic pain, post-operative pain, 15 migraine, arthralgia, neuropathies, nerve injury, diabetic neuropathy, neurodegeneration,

neurotic skin disorder, stroke, urinary bladder hypersensitivity, irritable bowel syndrome, a respiratory disorder such as asthma and chronic obstructive pulmonary diseases, irritation in skin, eye or mucous membrane, stomach-duodenal ulcer, inflammatory bowel disease, inflammatory disease, etc.

5

CLAIMS

1. A compound of formula I:



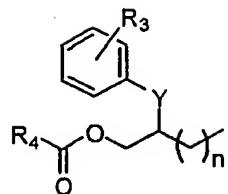
5 or a pharmaceutically acceptable salt thereof,

wherein,

X represents a sulfur atom or an oxygen atom;

R₁ represents a lower alkyl sulfonyl group having 1 to 5 carbon atoms, an aryl 10 sulfonyl group or a lower alkyl carbonyl group having 1 to 5 carbon atoms, which may be unsubstituted or substituted with halogen atom;

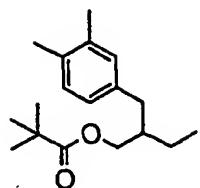
R₂ represents a hydrogen atom, a methoxy group or a halogen;



when A is -NHCH₂-, B represents

(wherein, n is 0 or

1), or when A is -CH₂-, B represents 4-t-butylbenzyl, 3,4-dimethylphenylethyl,



or an oleyl group;

Y represents -CH₂- or -CH₂CH₂-;

5 R₃ represents a hydrogen atom or a lower alkyl group having 1 to 5 carbon atoms; and

R₄ represents a lower alkyl group having 1 to 5 carbon atoms or phenyl group.

2. A pharmaceutical composition comprising the compound of formula I
10 according to claim 1 or a pharmaceutically acceptable salt thereof as an active ingredient together with a pharmaceutically acceptable carrier.

3. The pharmaceutical composition according to claim 2, wherein the

compound of formula I according to claim 1 or a pharmaceutically acceptable salt thereof as an active ingredient together with an acceptable carrier are present in an effective amount for preventing or treating pain, acute pain, chronic pain, neuropathic pain, post-operative pain, migraine, arthralgia, neuropathies, nerve injury, diabetic neuropathy, neurodegeneration, neurotic skin disorder, stroke, urinary bladder hypersensitivity, irritable bowel syndrome, a respiratory disorder such as asthma or chronic obstructive pulmonary disease, irritation of skin, eye or mucous membrane, fæces, stomach-duodenal ulcer, inflammatory bowel disease or inflammatory diseases.

10

4. A method for preventing or treating pain, acute pain, chronic pain, neuropathic pain, post-operative pain, migraine, arthralgia, neuropathies, nerve injury, diabetic neuropathy, neurodegeneration, neurotic skin disorder, stroke, urinary bladder hypersensitivity, irritable bowel syndrome, a respiratory disorder such as asthma or chronic obstructive pulmonary disease, irritation of skin, eye or mucous membrane, fæces, stomach-duodenal ulcer, inflammatory bowel disease or inflammatory diseases, wherein the method comprises administering a therapeutically effective amount of the compounds selected from the group consisting of compounds of formula I or pharmaceutically acceptable salts thereof.

5. The use of compounds selected from the group consisting of compounds of formula I or pharmaceutical acceptable salts thereof as antagonists of vanilloid receptors.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/KR01/01408

A. CLASSIFICATION OF SUBJECT MATTER

IPC7 C07C 33S/16, A61K 31/222

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C07C, A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
JournalElectronic data base consulted during the international search (name of data base and, where practicable, search terms used)
STN(REG, CAPLus) ; chemical structural search and caps? and vanilloid? and (thiourea or (thio (w) urea))

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	Jaewoo Lee et al, Thiourea analogues of resiniferatoxin as ligands for the vanilloid receptor, Bioorganic & Medicinal Chemistry Letters, Vol 5. No 13, pp 1331-1334, 1995, See the whole document	1-3, 5
A	US-A-6057451(2000.5.2) & US-A-6288091(2001.9.11) See the group 2(all schema), tables	1-3, 5
A	JP-A-58-55417(1983.4.1) & None See the whole document	1-3, 5
A	WO-A-99-37675(1999.7.29) & EP-A1-1047711(2000.11.2) & AU-A1-1998-72151(1998.1.22) See the whole document	1-3, 5
-----End of Documents-----		

 Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:
 "A" document defining the general state of the art which is not considered to be of particular relevance
 "E" earlier application or patent but published on or after the international filing date
 "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of citation or other special reason (as specified)
 "O" document referring to an oral disclosure, use, exhibition or other means
 "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
 "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
 "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
 "&" document member of the same patent family

Date of the actual completion of the international search
22 NOVEMBER 2001 (22.11.2001)

Date of mailing of the international search report
23 NOVEMBER 2001 (23.11.2001)

Name and mailing address of the ISA/KR
Korean Intellectual Property Office
Government Complex-Daejeon, Dunsan-dong, Seo-gu, Daejeon
Metropolitan City 302-701, Republic of Korea
Facsimile No. 82-42-472-7140

Authorized officer
PARK, Kil Chae
Telephone No. 82-42-481-5536



INTERNATIONAL SEARCH REPORT

International application No.

PCT/KR01/01408

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: 4
because they relate to subject matter not required to be searched by this Authority, namely:
Method for treatment of the human body by therapy.
2. Claims Nos.:
because they relate to part of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Search Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be established without effort justifying an additional fee, this Authority did not invite payment of any addition fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.